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? s rituximab  
S1 36 RITUXIMAB  
? s atcc  
S2 10152 ATCC  
? s s1 and s2  
36 S1  
10152 S2  
S3 2 S1 AND S2  
? t s3/3,k,ab/1-2

3/3,K,AB/1  
DIALOG(R) File 340: CLAIMS(R) /US Patent  
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Dialog Acc No: 10069121 IFI Acc No: 2002-0012665 IFI Acc No: 2002-0003424

Document Type: C

COMBINED USE OF ANTI-CYTOKINE ANTIBODIES OR ANTAGONISTS AND ANTI-CD20 FOR TREATMENT OF B CELL LYMPHOMA; ADMINISTERING AN ANTI-CYTOKINE ANTIBODY OR FRAGMENT THEREOF OR CYTOKINE ANTAGONIST TO A PATIENT DIAGNOSED WITH A HEMATOLOGIC MALIGNANCY OR A SOLID, NON-HEMATOLOGIC TUMOR PRIOR, CONCURRENT OR AFTER ADMINISTRATION CHEMOTHERAPEUTIC AGENT

Inventors: Hanna Nabil (US)

Assignee: Unassigned Or Assigned To Individual

Assignee Code: 68000

Publication (No,Date), Applic (No,Date):

US 20020012665 20020131 US 2001822672 20010402

Publication Kind: A1

Priority Applic(No,Date): US 2001822672 20010402

Provisional Applic(No,Date): US 60-193467 20000331

Abstract: The present invention discloses combined therapies for treating hematologic malignancies, including B cell lymphomas and leukemias or solid non-hematologic tumors, comprising administration of anti-cytokine antibodies or antagonists to inhibit the activity of cytokines which play a role in perpetuating the activation of B cells. The administration of such antibodies and antagonists, particularly anti-IL10 antibodies and antagonists, is particularly useful for avoiding or decreasing the resistance of hematologic malignant cells or solid tumor cells to chemotherapeutic agents and anti-CD20 or anti-CD22 antibodies. The invention also provides combination therapies for solid tumors having B cell involvement comprising the administration of an anti-cytokine antibody and a B cell depleting antibody such as RITUXAN registered .

Non-exemplary Claims: ...41. The method of claim 40, where said chimeric anti-CD20 antibody is **Rituximab** registered42. The method of claim 41, wherein said **Rituximab** registered is administered at a dosage of 0.4 to 20 mg/kg body weight...wherein said anti-CD20 antibody is **Rituxan** registered , a chimeric anti-CD20 antibody produced by **ATCC** 69119...71. The method of claim 70 wherein said antibody is **Rituxan** registered produced by **ATCC** 69119... .

3/3,K,AB/2  
DIALOG(R) File 340: CLAIMS(R) /US Patent  
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Dialog Acc No: 3887549 IFI Acc No: 0020642

Document Type: C

METHODS FOR THE PREVENTION AND TREATMENT OF CANCER USING ANTI-C3B(I) ANTIBODIES; CLASSIC COMPLEMENT PATHWAY, C3 CONVERTASE

Inventors: Chung Leland (US); Nardin Alessandra (FR); Sokoloff Mitchell H (US); Sutherland William M (US); Taylor Ronald (US)

Assignee: Virginia, University of Alumni Patents Foundation  
Assignee Code: 22061  
Publication (No,Date), Applic (No,Date):  
US 6572856 20030603 US 2000724620 20001128  
Publication Kind: B  
Cont.-in-part Pub(No),Applic(No,Date): US  
88392500 19880209  
Priority Applic(No,Date): US 2000724620 20001128; US 88392500 19880209

Abstract: The present invention relates to the treatment and prevention of cancer, viral infections and microbial infections by the administration of anti-C3b(i) antibodies. The present invention also relates to methods of treating and preventing cancer, viral infection, or microbial infection in an animal comprising administering to said animal IgG antibodies, IgM antibodies and/ or complement components in combination with antibodies specific for C3b(i). The present invention also relates methods of treating and preventing cancer, viral infection or microbial infection in an animal comprising administrating said animal antibodies that immunospecifically bind to one or more cancer cell antigens, viral antigens or microbial antigens, respectively, in combination with antibodies immunospecific for C3b(i). The present invention further relates to the detection, imaging, diagnosis and monitoring of cancer utilizing C3b(i) specific antibodies.

Non-exemplary Claims: ...of the anti-C3b(i) antibodies is 3E7 produced by the hybridoma deposited with the **ATCC** as Accession No. PTA-4090...

...wherein the anti-C3b(i) antibody is 3E7 produced by the hybridoma deposited with the **ATCC** as Accession No. PTA-4090...

...41. The method of claim 24, wherein the anti-CD20 antibody is **rituximab**.

...

...wherein the anti-C3b(i) antibody is 3E7 produced by the hybridoma deposited with the **ATCC** as Accession No. PTA-4090 and the anti-CD20 antibody is **rituximab**.

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Feb-27-2004 10:46

From-PILLSBURY WINTHROP

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T-207 P.001/001 F-876

U.S. Application No. 09/762,587  
Attorney Ref. No. 037003-0277847

Examiner Davis:

Pages 2-3 of our previous facsimile correspondence (the Bexxar product insert) can be found at:

<http://www.corixa.com/Bexxar/BexxarPackageInsert.pdf>

The complete citation for the Coleman et al. meeting abstract is as follows:

Coleman M, Kaminski MS, Knox SJ, Zelenetz AD, et al. The BEXXAR Therapeutic Regimen (Tositumomab and Iodine 131 Tositumomab) Produced Durable Complete Remissions in Heavily Pretreated Patients with Non-Hodgkin's Lymphoma (NHL), Rituximab-Relapsed/Refractory Disease, and Rituximab-Naive Disease. *Proc Am Soc Hem. Blood.* 2003; 102(11):29a, Abstract #89.

The abstract was presented as an oral session at the 45th annual meeting of the American Society of Hematology on December 7, 2003.

*Julie Bronder Meigs*